**DIPHEN- benzocaine, benzalkonium chloride, lidocaine hydrochloride, hydrocortisone, bacitracin zinc, neomycin sulfate, polymyxin b sulfate, calcium carbonate, ibuprofen, loratadine, acetaminophen, dextromethorphan hydrobromide, guaifenesin, phenylephrine hydrochloride, diphenhydramine hydrochloride, potassium chloride, magnesium oxide, meclizine hydrochloride, and bismuth subsalicylate**
**Remedy Pack LLC**

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**Diphen**

**REMEDY PACK**

**Burn Cream**

***Drug Facts***

## INGREDIENTS

### Active ingredients

Benzalkonium Chloride 0.13%, Lidocaine HCl 0.5%

### Purpose

Topical antiseptic, Topicall analgesic

## USES

First aid to help prevent infection in minor cuts, scrapes and burns.

For the temporary relief of pain and itching associated with:

- sunburn
- insect bites
- cuts
- minor skin irritations
- scrapes
- minor burns

## WARNING

For external use only.

### Do not use

- in the eyes
- over large areas of the body or on deep puncture wounds, animal bites, or serious burns
- in large quantities, particularly over raw surfaces or blistered areas

### Stop use and ask doctor if

- the condition gets worse
- condition clears up and recurs within a few days
- condition persists for more than 7 days

If pregnant or breast feeding, ask a health care professional before use.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

## DIRECTIONS

Adults and children 2 years and over:
- clean the affected area
- apply a small amount of this product on the area 3 to 4 times daily
- may be covered with a sterile bandage
- ***Children under 12 years:*** consult a doctor
- ***Children under 2 years:*** consult a doctor

## OTHER INFORMATION
- store in a cool, dry area 59° to 79°F (15° to 25°C)
- tamper evident sealed packets
- do not use any opened or torn packets

## INACTIVE INGREDIENTS

decolorized aloe vera, emulsifying wax, ethyl alcohol, methylparaben, mineral oil, paraffin, propylparaben, purified water, white petrolatum, white wax

**Triple Antibiotic**

***Drug Facts***

## INGREDIENTS

### Active Ingredient (in each gram)

Bacitracin zinc (400 units), Neomycin sulfate 5 mg (equivalent to 3.5 mg of Neomycin), Polymyxin-B sulfate 5000 units

### Purposes

First aid antibiotics

## USES

First aid to help prevent infection in:
- minor cuts
- scrapes

- burns

**WARNING**
- For external use only

**Do not use**
- in the eyes
- over large areas of the body if you are allergic to any of the ingredients longer than 1 week unless directed by a doctor

**Ask a doctor before use**

in case of deep or puncture wounds, animal bites, or serious burns

**Stop use and ask a doctor if**

the condition persists or gets worse a rash or other allergic reaction develops

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center immediately.

**DIRECTIONS**

clean the affected area apply a small amount of this product (an amount equal to the surface area of the tip of a finger) on the area 1 to 3 times daily
may be covered with a sterile bandage

**OTHER INFORMATION**
- store at room temperature 15°C to 30°C (59°F to 86°F) (do not freeze) tamper evident. Do not use if packet is torn, cut or opened.
- avoid excessive heat and humidity

**INACTIVE INGREDIENTS**

mineral oil, white petrolatum

**Hydrocortisone Cream**

***Drug Facts***

**INGREDIENTS**

**Active Ingredient (in each gram)**

Hydrocortisone 1.0%

**Purpose**

Anti-Itch

**USES**

- eczema
- insect bites
- poison ivy
- Poison oak
- poison sumac
- Cosmetics
- jewelry
- soaps detergents
- seborrheic
- dermatitis
- Psoriasas

Other uses of this product should only be under the advice and supervision of a doctor.

**WARNING**

- For external use only
- avoid contact with the eyes
- if condition worsens, or symptoms persist for more than 7 days or clear up and occur again within a few days, stop use of this product and do not begin use of any other hydrocortisone product unless you have consulted a doctor
- do not use for the treatment of diaper rash.
- Keep out of reach of children.
- If swallowed, get medical help or consult a poison control center right away.

**DIRECTIONS**

***Adult and children (2 years and over):*** apply to affected not more than 3 to 4 times daily

***Children under 2 years:*** Consult a doctor.

**OTHER INFORMATION**

- store at room temperature 59-86ºF (15-30ºC)
- do not freeze
- do not use any opened or torn packets.

**INACTIVE INGREDIENTS**

emulsifying wax, ethanol, methylparaben, mineral oil, paraffin, petrolatum, propylparaben, purified water, white Wax.

**Oral Pain Relief Gel**

***Drug Facts***

## INGREDIENTS

### Active ingredients

Benzocaine 20%

### Purpose

Oral Anesthetic

## USES

For oral mucosal use only, as directed by dentist. For the temporary relief of pain due to minor dental procedures.

## METHEMOGLOBINEMIA WARNING

Use of this product may cause methemoglobinemia, a serious condition that must be treated promptly because it reduces the amount of oxygen carried in blood.

This can occur even if you have used this product before. Stop use and seek immediate medical attention if you or a child in your care develops:

- pale, gray, or blue colored skin (cyanosis)
- headache
- rapid heart rate
- shortness of breath
- dizziness or lightheadedness
- fatigue or lack of energy

## ALLERGY ALERT

do not use this product if you have a history of allergy to local anesthetics such as procaine, butacaine, benzocaine, or other " caine" anesthetics.

### Do not use

For more than 7 days unless directed by a physician. If sore mouth symptoms do not improve in 7 days; irritation, pain, or redness persists or worsens; or if swelling, rash or fever develops, see your physician promptly.

- for teething
- in children under 2 years of age

### When using this product

avoid contact with eyes. If it occurs, flush with water.

***Do not exceed recommended dosage.***

- If more than used for pain is accidently swallowed, get medical help or contact a Poison Control Center right away.

- If pregnant or breast feeding,ask a health care professional before use.

- Keep out of reach of children.

**DIRECTIONS**
- Apply only amount needed to the oral mucosa to prevent or relieve pain.
- children under 2 years of age: do not use

**OTHER INFORMATION**
- store at room temperature 15°C to 30°C (59°F to 86°F)
- protect from freezing

**INACTIVE INGREDIENTS**

Flavoring, PEG 3350, PEG 400, sodium sacchrin, water.

May contain blue #1, green #3, green #5 , red #3, red #28, red #40, yellow #5, (tartrazine), yellow #6, as color additive.

**Alcalak**

***Drug Facts***

**INGREDIENTS**

**Active ingredient (in each tablet)**

Calcium Carbonate 420mg

**Purpose**

Antacid

**INACTIVE INGREDIENTS[1]**

aspartame[1], croscarmellose sodium[1], gum acacia[1], magnesium stearate, maltodextrin, mineral oil[1], mint flavor, sorbitol[1], sucrose[1]

---

1 may contain

**USES**

For the relief of the following symptoms associated with
- acid indigestion
- sour stomach
- heartburn
- upset stomach

**WARNING**

**Do not use**

- the maximum dosage of this product for more than 2 weeks, except under the advice and supervision of a physician, or take more than 19 tablets in a 24 hour period.

**Ask a doctor or pharmacist before use if you are**

- presently taking a prescription drug. Antacids may interact with certain prescription drugs.

**Stop use and ask a doctor if**

- symptoms last more than 2 weeks

If pregnant or breastfeeding, ask a health professional before use.

Keep out of reach of children.

**DIRECTIONS**

- do not use more than directed

***Adults and children:*** (12 years and older) Chew 2 tablets every 2 or 3 hours as symptoms occur or as directed by a physician. Do not exceed 19 tablets in 24 hours.

***Children under 12 years:*** Do not give to children under 12 years of age.

**OTHER INFORMATION**

- Phenylketonurics: contains phenylalanine 1.5mg per tablet
- each tablet contains 168mg of elemental calcium
- store at room temperature 59º-86ºF (15º-30ºC) in a dry place
- tamper-evident sealed packets
- do not use any opened or torn packets

**Cold Relief**

***Drug Facts***

**INGREDIENTS**

| | |
|---|---|
| **Active ingredient (in each tablet):** Acetaminophen 325mg | **Purpose:** Pain reliever/ fever reducer |
| **Active ingredient (in each tablet):** Dextromethorphan Hydrobromide 15mg | **Purpose:** Cough suppressant |
| **Active ingredient (in each tablet):** Guaifenesin 200mg | **Purpose:** Expectorant |
| **Active ingredient (in each tablet):** Phenylephrine HCl 5mg | **Purpose:** Nasal decongestant |

**INACTIVE INGREDIENTS***

maltodextrin, microcrystalline cellulose, povidone, sodium starch glycolate, starch, stearic acid

**USES**

Temporarily relieves these symptoms due to hay fever or other upper respiratory allergies

- cough
- sore throat
- minor aches and pains
- headache
- nasal congestion
- helps loosen phlegm (mucus) and thin bronchial secretions to make coughs more productive

Temporarily reduces fever.

**WARNING**

**Liver warning**

This product contains acetaminophen. Severe liver damage may occur if you take:

- more than 4,000mg of acetaminophen in 24 hours
- with other drugs containing acetaminophen
- 3 or more alcoholic drinks every day while using this product

**Allergy alert**

Acetaminophen may cause severe skin reactions. Symptoms may include:

- skin reddening
- blisters
- rash

If a skin reaction occurs, stop use and seek medical help right away.

**Do not use**

- with any other drug containing acetaminophen (prescription or nonprescription). If you are not sure whether a drug contains acetaminophen, ask a doctor or pharmacist.
- if you have ever had an allergic reaction to this product or any of the ingredients.
- if you are now taking a prescription monoamine oxidase inhibitor (MAOI) (certain drugs for depression, psychiatric or emotional conditions, or Parkinson's disease), or for 2 weeks after stopping the MAOI drug. If you do not know if your prescription drug contains an MAOI, ask a doctor or pharmacist before taking this product.

**Ask a doctor before use if you have**

- liver disease
- heart disease

- high blood pressure
- thyroid disease
- diabetes
- trouble urinating due to an enlarged prostate gland
- cough that occurs with too much phlegm (mucus)
- persistent or chronic cough that lasts as occurs with smoking, asthma, chronic bronchitis or emphysema

**Ask a doctor or pharmacist before use if you are**
- taking the blood thinning drug warfarin

**When using this product**
- do not use more than directed

**Stop use and ask a doctor if**
- new symptoms occur
- redness or swelling is present
- pain or nasal congestion gets worse or lasts for more than 7 days
- fever gets worse or lasts for more than 3 days
- you get nervous, dizzy or sleepless
- cough comes back or occurs with rash or headache that lasts.

These could be signs of a serious condition.

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away (1-800-222-1222).

Prompt medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

**DIRECTIONS**

***Adults and children:*** (12 years and older) Take 2 tablets with water every 6- 8 hours as needed. Do not take more than 8 tablets in 24 hours.

***Children under 12 years:*** Do not give to children under 12 years of age.

**OTHER INFORMATION**
- store at room temperature 59º-86ºF (15º-30ºC)
- avoid excessive heat and humidity
- tamper-evident sealed packets
- do not use any opened or torn packets

**Medi-Meclizine**

***Drug Facts***

**INGREDIENTS**

**Active ingredient (in each tablet)**

Meclizine Hydrochloride 25mg

**Purpose**

Antiemetic

**INACTIVE INGREDIENTS***

anhydrous lactose, colloidal silicon dioxide, corn starch, D&C yellow #10, magnesium stearate, microcrystalline cellulose, sodium starch glycolate

**USES**

For the prevention and treatment of nausea, vomiting, or dizziness associated with motion sickness.

For the reduction of fever.

**WARNING**

**Do not use**

- for children under 12 years of age unless directed by a doctor
- for frequent or prolonged use except under the advice of a doctor

**Ask a doctor before use if you have**

- breathing problems such as emphysema or chronic bronchitis
- glaucoma
- difficulty in urination due to enlargement of the prostate gland if you are
- taking sedatives or tranquilizers

**When using this product**

- drowsiness may occur
- alcohol, sedatives and tranquilizers may increase the drowsiness effect
- avoid alcoholic beverages while taking this product
- use caution when driving a motor vehicle or operating machinery

Do not exceed recommended dosage.

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away.

**DIRECTIONS**

- do not use more than directed
- to prevent motion sickness, take the first dose one hour before starting activity

***Adults and children:*** (12 years and older) 1 to 2 tablets once daily or as directed by a doctor. Do not exceed 2 tablets in 24 hours.

***Children under 12 years:*** Do not give to children under 12 years of age.

**OTHER INFORMATION**

- store at room temperature 59- 86º F (15-30ºC)
- tamper-evident sealed packets
- do not use any opened or torn packets

**I-Prin**

***Drug Facts***

**INGREDIENTS**

**Active ingredient (in each tablet)**

Ibuprofen (NSAID[2]) 200mg

---

2 nonsteroidal anti-inflammatory drug

**Purpose**

Pain reliever/fever reducer

**INACTIVE INGREDIENTS[3]**

carnauba wax[3], corn starch, hypromellose[3], iron oxide red, lactose[3], magnesium stearate[3], microcrystalline cellulose[3], polydextrose[3], polyethylene glycol, polyvinyl alcohol[3], povidone (K-30)[3], silicon dioxide, sodium starch glycolate, stearic acid, talc[3], titanium dioxide

---

3 may contain

**USES**

Temporarily relieves minor aches and pains associated with

- headache
- toothache
- backache
- menstrual cramps
- common cold
- muscular aches
- minor arthritis pain

Temporarily reduces fever.

## WARNING

### Allergy alert

Ibuprofen may cause a severe allergic reaction, especially in people allergic to aspirin.

**Symptoms may include:**

- hives
- skin reddening
- facial swelling
- rash
- asthma (wheezing)
- blisters
- shock

If an allergic reaction occurs, stop use and seek medical help right away.

### Stomach bleeding warning

This product contains an NSAID, which may cause severe stomach bleeding.

**The chance is higher if you:**

- are age 60 or older
- have had stomach ulcers or bleeding problems
- take a blood thinning (anticoagulant) or steroid drug
- take other drugs containing prescription or nonprescription NSAIDs (aspirin, ibuprofen, naproxen, or others)
- have 3 or more alcoholic drinks every day while using this product
- take more or for a longer time than directed

Heart attack or stroke warning: NSAIDS, except aspirin, increase the risk of heart attack, heart failure, and stroke.

These can be fatal. The risk is higher if you use more than directed or for longer than directed.

### Do not use

- if you have ever had an allergic reaction to any other pain reliever/ fever reducer
- right before or after heart surgery

### Ask a doctor before use if

- you have problems or serious side effects from taking pain relievers or fever reducers
- stomach bleeding warning applies to you
- you have a history of stomach problems such as heartburn
- you have high blood pressure, heart disease, liver cirrhosis, kidney disease, asthma or had a stroke
- you are taking a diuretic

### Ask a doctor or pharmacist before use if you are

- taking aspirin for heart attack or stroke, because ibuprofen may decrease this benefit

of aspirin
- under a doctor's care for any serious condition
- taking any other drug

**When using this product**
- take with food or milk if stomach upset occurs

**Stop use and ask a doctor if**
- you experience any of the following signs of stomach bleeding:
    - feel faint
    - vomit blood
    - have bloody or black stools
    - have stomach pain that does not get better
- you have symptoms of heart problems or stroke
    - chest pain
    - trouble breathing
    - weakness in one part or side of body
    - slurred speech
    - leg swelling
- pain gets worse or lasts more than 10 days
- fever gets worse or lasts more than 3 days
- redness or swelling is present in the painful area
- any new or unexpected symptoms occur

If pregnant or breast-feeding, ask a health professional before use. It is especially important not to use ibuprofen during the last 3 months of pregnancy unless specifically directed to do so by a doctor because it may cause problems in the unborn child or complications during delivery.

Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away.

**DIRECTIONS**
- do not take more than directed
- the smallest effective dose should be used
- do not take longer than 10 days, unless directed by a doctor (see Warnings)

***Adults and children:*** (12 years and older) Take 1 tablet every 4 to 6 hours while symptoms persist. If pain or fever does not respond to 1 tablet, 2 tablets may be used. Do not exceed 6 tablets in 24 hours, unless directed by a doctor.

***Children under 12 years:*** Do not give to children under 12 years of age.

**OTHER INFORMATION**
- read all product information before using
- store at 68-77°F (20-25°C)

- avoid excessive heat 104°F (above 40°C)
- tamper-evident sealed packets
- do not use any opened or torn packets

**Diphen**

***Drug Facts***

**INGREDIENTS**

**Active ingredient (in each tablet)**

Diphenhydramine HCl 25mg

**Purpose**

Antihistamine

**INACTIVE INGREDIENTS[4]**

carnauba wax[4], colloidal silicon dioxide, croscarmellose sodium, D&C red #27, dicalcium phosphate[4], hypromellose, lactose[4], magnesium stearate, microcrystalline cellulose, polyethylene glycol, polysorbate[4], titanium dioxide

---

4 may contain

**USES**

**Active ingredients**

Diphenhydramine HCl 25mg

**Purpose**

Antihistamine

**Inactive Ingredients[5]**

carnauba wax[5], colloidal silicon dioxide, croscarmellose sodium, D&C red #27, dicalcium phosphate[5], hypromellose, lactose[5], magnesium stearate, microcrystalline cellulose, polyethylene glycol, polysorbate[5], titanium dioxide

---

5 may contain

**WARNING**

**Do not use**
- to make a child sleepy
- with any other product containing diphenhydramine, even one that is used on skin

**Ask a doctor before use if you have**
- a breathing problem such as emphysema or chronic bronchitis
- difficulty in urination due to enlargement of the prostate gland
- glaucoma

**Ask a doctor or pharmacist before use if you are**
- taking sedatives or tranquilizers

**When using this product**
- marked drowsiness may occur
- avoid alcohol beverages
- alcohol, sedatives and tranquilizers may increase the drowsiness effect
- use caution when driving a motor vehicle or operating machinery
- excitability may occur, especially in children

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children. In case of overdose, contact a physician or Poison Control Center immediately.

**DIRECTIONS**
- do not use more than directed

***Adults and children:*** (12 years and older) Take 1 to 2 caplets every 4 to 6 hours as needed. Do not take more than 12 caplets in 24 hours, or as directed by a doctor.

***Children under 12 years:*** Do not give to children under 12 years of age.

**OTHER INFORMATION**
- each caplet may contain: calcium 25mg
- protect from light
- use by expiration date on packet
- store at room temperature 59º-86ºF (15º-30ºC)
- tamper-evident sealed packets
- do not use any opened or torn packets

**Extra Strength APAP**

***Drug Facts***

**INGREDIENTS**

**Active ingredient (in each tablet)**

Acetaminophen 500mg

**Purpose**

Pain reliever/fever reducer

## INACTIVE INGREDIENTS[6]

corn starch, hypromellose, maltodextrin[6], microcrystalline cellulose[6], polyethylene glycol, povidone[6], pregelatinized starch[6], sodium starch glycolate[6], stearic acid, titanium dioxide[6]

---

6 may contain

## USES

For the temporary relief of minor aches and pains associated with

- headache
- muscular aches
- minor arthritis pain
- common cold
- toothache
- menstrual cramps

For the reduction of fever.

## WARNING

### Liver warning

This product contains acetaminophen. Severe liver damage may occur if you take:

- more than 8 tablets in 24 hours, which is the maximum daily amount
- with other drugs containing acetaminophen
- 3 or more alcoholic drinks every day while using this product

### Allergy alert

Acetaminophen may cause severe skin reactions.

**Symptoms may include:**

- skin reddening
- blisters
- rash

If a skin reaction occurs, stop use and seek medical help right away.

### Do not use

- with any other drug containing acetaminophen (prescription or nonprescription). If you are not sure whether a drug contains acetaminophen, ask a doctor or pharmacist.
- for more than 10 days for pain unless directed by a doctor
- for more than 3 days for fever unless directed by a doctor

### Ask a doctor before use if you have

- liver disease

**Ask a doctor or pharmacist before use if**
- you are taking the blood thinning drug warfarin

**Stop using and ask a doctor if**
- symptoms do not improve
- new symptoms occur
- pain or fever persists or gets worse
- redness or swelling is present

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children. In case of accidental overdose, get medical help or contact a Poison Control Center right away. Prompt medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

**DIRECTIONS**
- do not use more than directed

***Adults and children:*** (12 years and older) Take 2 tablets every 4 to 6 hours as needed. Do not take more than 8 tablets in 24 hours.

***Children under 12 years:*** Do not give this adult strength product to children under 12 years of age; this will provide more than the recommended dose (overdose) and may cause liver damage.

**OTHER INFORMATION**
- store at room temperature 59º-86ºF (15º-30ºC)
- tamper-evident sealed packets
- do not use any opened or torn packets

**Loradamed**

***Drug Facts***

**INGREDIENTS**

**Active ingredient (in each tablet)**

Loratadine 10mg

**Purpose**

Antihistamine

**INACTIVE INGREDIENTS***

corn starch, lactose monohydrate, magnesium stearate, pregelatinized starch

## USES

Temporarily relieves these symptoms due to hay fever or other upper respiratory allergies

- runny nose
- itchy, watery eyes
- sneezing
- itching of the nose or throat

## WARNING

Do not use if you have ever had an allergic reaction to this product or any of its ingredients.

Ask a doctor before use if you have liver or kidney disease. Your doctor should determine if you need a different dose.

When using this product do not take more than directed. Taking more than directed may cause drowsiness.

### Stop use and ask a doctor if

- *an allergic reaction to this product occurs. Seek medical help right away.*

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

## DIRECTIONS

***Adults and children:*** (12 years and older) Take 1 tablet daily; not more than 1 tablet in 24 hours.

**Children under 12 years:** Do not give to children under 12 years of age.

***Consumers with liver or kidney disease:*** Ask a doctor before using.

## OTHER INFORMATION

- store at room temperature 68º-77ºF (20º-25ºC)
- protect from excessive moisture
- tamper-evident sealed packets
- do not use any opened or torn packets

### Medi-Lyte

***Drug Facts***

## INFORMATION

Serving Size: 2 tablets

Servings Per Packet: 1

**OTHER INGREDIENTS**

microcrystalline cellulose, silicon dioxide, stearic acid, magnesium stearate

| ***Amount Per Serving*** | ***% Daily Value*** |
|---|---|
| Calcium (from 27.0 mg calcium carbonate) 10.8 mg | 1.06% |
| Potassium (from 80 mg potassium chloride) 40mg | 1.15% |
| Magnesium (from 20 mg magnesium oxide) 12 mg | 3.0% |
| Carbohydrates 6 mg | >1% |
| Calories 1.5 | >1% |
| Protein | 0% |
| Fat | 0% |

**USES**

Nutritional support for the following symptoms due to excessive loss of perspiration

- heat fatigue
- muscle cramps
- heat exhaustion
- heat stroke
- replaces lost electrolytes
- helps provide rapid rehydration

*This statement has not been evaluated by the Food and Drug administration. This product is not intended to diagnose, treat, cure, or prevent any disease.

**WARNING**

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away.

**DIRECTIONS**

***Adults and children:*** (12 years and older)

- take 2 tablets as needed with a full glass of water
- may be repeated every hour as needed
- do not exceed 20 tablets in 24 hours

***Children under 12 years:***

- Do not give to children under 12 years of age

**OTHER INFORMATION**

- store at room temperature 59º-86º F (15º-30º C)
- avoid excessive heat and humidity
- tamper-evident sealed packets
- do not use any opened or torn packets
- no sodium added

**Diotame**

***Drug Facts***

**INGREDIENTS**

**Active ingredient (in each tablet)**

Bismuth Subsalicylate 262mg (each tablet contains 102mg salicylate)

**Purpose**

Upset stomach reliever/antidiarrheal

**INACTIVE INGREDIENTS***

acacia gum, aspartame, calcium carbonate, D&C red #27, dextrates, flavoring, magnesium stearate, maltodextrin, microcrystalline cellulose, silicon dioxide

**USES**

Temporarily relieves

- travelers' diarrhea
- diarrhea
- upset stomach due to overindulgence in food and drink, including
    - heartburn
    - indigestion
    - nausea
    - gas
    - belching
    - fullness

**WARNING**

**Reye's syndrome**

Children and teenagers who have or are recovering from chicken pox or flu-like symptoms should not use this product. When using this product, if changes in behavior with nausea and vomiting occur, consult a doctor because these symptoms could be an early sign of Reye's Syndrome a rare but serious illness.

**Allergy alert**

**Contains salicylate. Do not take if you are:**
- allergic to salicylates (including aspirin)
- taking other salicylate products

**Do not use if you have**
- bloody or black stool
- an ulcer
- a bleeding problem

**Ask a doctor before use if you have**
- fever
- mucus in the stool

**Ask a doctor or pharmacist if you are taking any drug for**
- anticoagulation (thinning of the blood)
- diabetes
- gout
- arthritis

**Stop use and ask a doctor if**
- symptoms get worse
- ringing in the ears or loss of hearing occurs
- diarrhea lasts more than 2 days

When using this product a temporary and harmless darkening of the tongue and/or stool may occur. Stool darkening should not be confused with melena.

If pregnant or breast-feeding, ask a health professional before use.

Keep this and all drugs out of reach of children. In case of accidental overdose, contact a physician or poison control center immediately.

**DIRECTIONS**
- do not use more than directed
- chew or crush tablets completely before swallowing
- do not swallow tablets whole
- use until diarrhea stops but not more than 2 days
- drink plenty of clear fluids to help prevent dehydration, which may accompany diarrhea
- do not exceed 16 tablets in 24 hours

***Adults and children:*** (12 years and older) Chew 2 tablets every 1/2 to 1 hour or 4 tablets every hour as needed.

***Children under 12 years:*** Do not give to children under 12 years of age.

**OTHER INFORMATION**

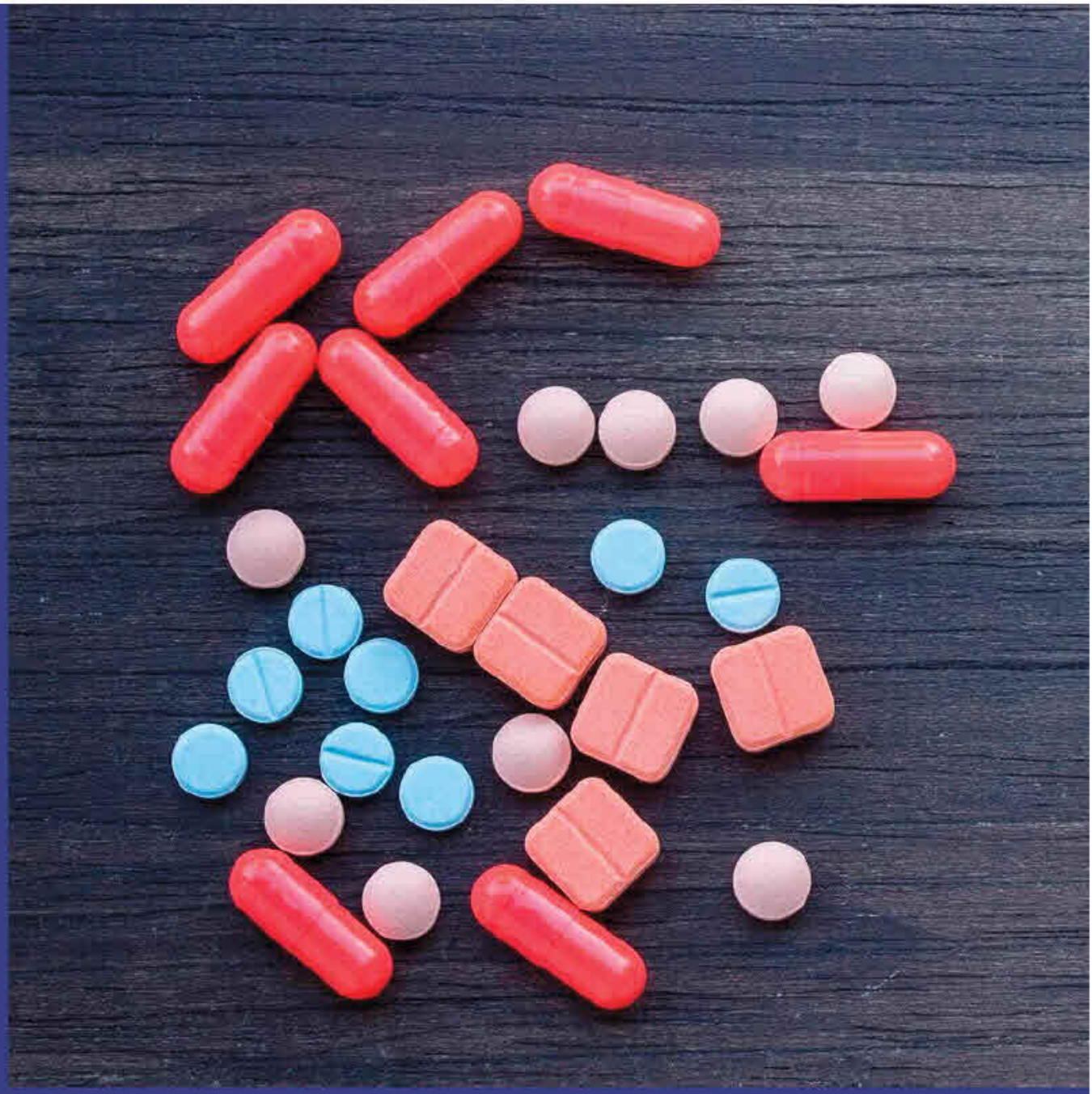
- Phenylketonurics: contains phenylalanine 1.1mg per tablet
- each tablet contains 73mg of elemental calcium
- store at room temperature 59º-86ºF (15º-30ºC)
- tamper-evident sealed packets
- do not use any opened or torn packets

**PRINCIPAL DISPLAY PANEL - Kit Carton**

REMEDY PACK

DRUG FACTS

www.theremedypack.com



# REMEDY PACK

# DRUG FACTS

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REMEDY PACK

## Triple Antibiotic

### Drug Facts

***INGREDIENTS***
**Active Ingredient (in each gram):** Bacitracin zinc (400 units), Neomycin sulfate 5 mg (equivalent to 3.5 mg of Neomycin), Polymyxin-B sulfate 5000 units
**Purposes:** First aid antibiotics

***USES***
First aid to help prevent infection in: • minor cuts • scrapes • burns

***WARNING***
• For external use only
***Do not use***
• in the eyes • over large areas of the body if you are allergic to any of the ingredients longer than 1 week unless directed by a doctor

***Ask a doctor before use:*** in case of deep or puncture wounds, animal bites, or serious burns

***Stop use and ask a doctor if:*** the condition persists or gets worse a rash or other allergic reaction develops

Keep out of reach of children.
If swallowed, get medical help or contact a Poison Control Center immediately.

***DIRECTIONS***
clean the affected area apply a small amount of this product (an amount equal to the surface area of the tip of a finger) on the area 1 to 3 times daily
may be covered with a sterile bandage

***OTHER INFORMATION***
• store at room temperature 15°C to 30°C (59°F to 86°F) (do not freeze) tamper evident. Do not use if packet is torn, cut or opened.
• avoid excessive heat and humidity

***INACTIVE INGREDIENTS***
mineral oil, white petrolatum

## Oral Pain Relief Gel

### Drug Facts

***INGREDIENTS***
**Active Ingredients:** Benzocaine 20% **Purpose:** Oral Anesthetic

***USES***
For oral mucosal use only, as directed by dentist. For the temporary relief of pain due to minor dental procedures.

***METHEMOGLOBINEMIA WARNING:***
Use of this product may cause methemoglobinemia, a serious condition that must be treated promptly because it reduces the amount of oxygen carried in blood. This can occur even if you have used this product before. Stop use and seek immediate medical attention if you or a child in your care develops:
• pale, gray, or blue colored skin (cyanosis) • headache • rapid heart rate • shortness of breath • dizziness or lightheadedness • fatigue or lack of energy

***ALLERGY ALERT***
do not use this product if you have a history of allergy to local anesthetics such as procaine, butacaine, benzocaine, or other " caine" anesthetics.

***Do not use***
For more than 7 days unless directed by a physician. If sore mouth symptoms do not improve in 7 days; irritation, pain, or redness persists or worsens; or if swelling, rash or fever develops, see your physician promptly.
• for teething • in children under 2 years of age

***When using this product:*** avoid contact with eyes. If it occurs, flush with water.

***Do not exceed recommended dosage.***
- If more than used for pain is accidently swallowed, get medical help or contact a Poison Control Center right away.
- If pregnant or breast feeding,ask a health care professional before use.
- Keep out of reach of children.

***DIRECTIONS***
- Apply only amount needed to the oral mucosa to prevent or relieve pain.
- children under 2 years of age: do not use

***OTHER INFORMATION***
- store at room temperature 15°C to 30°C (59°F to 86°F)
- protect from freezing

***INACTIVE INGREDIENTS***
Flavoring, PEG 3350, PEG 400, sodium saccharin, water.
May contain blue #1, green #3, green #5 , red #3, red #28, red #40, yellow #5, (tartrazine), yellow #6, as color additive.

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## Burn Cream

### *Drug Facts*

***INGREDIENTS***
**Active ingredients:** Benzalkonium Chloride 0.13%, Lidocaine HCl 0.5% **Purpose:** Topical antiseptic, Topicall analgesic

***USES***
First aid to help prevent infection in minor cuts, scrapes and burns.
For the temporary relief of pain and itching associated with: • sunburn • insect bites • cuts • minor skin irritations • scrapes • minor burns

***WARNING***
For external use only.

***Do not use*** • in the eyes • over large areas of the body or on deep puncture wounds, animal bites, or serious burns
• in large quantities, particularly over raw surfaces or blistered areas

***Stop use and ask doctor if*** • the condition gets worse • condition clears up and recurs within a few days • condition persists for more than 7 days
If pregnant or breast feeding, ask a health care professional before use.
Keep out of reach of children.
If swallowed, get medical help or contact a Poison Control Center right away.

***DIRECTIONS***
Adults and children 2 years and over:
• clean the affected area • apply a small amount of this product on the area 3 to 4 times daily • may be covered with a sterile bandage

• ***Children under 12 years:*** consult a doctor • ***Children under 2 years:*** consult a doctor

***OTHER INFORMATION***
• store in a cool, dry area 59° to 79°F (15° to 25°C) • tamper evident sealed packets • do not use any opened or torn packets

***INACTIVE INGREDIENTS***
decolorized aloe vera, emulsifying wax, ethyl alcohol, methylparaben, mineral oil, paraffin, propylparaben, purified water, white petrolatum, white wax

# Hydrocortisone Cream

## *Drug Facts*

***INGREDIENTS***
**Active Ingredient (in each gram):** Hydrocortisone 1.0% **Purpose:** Anti-Itch

***USES***
• eczema • insect bites • poison ivy • Poison oak • poison sumac • Cosmetics • jewelry • soaps detergents • seborrheic • dermatitis • Psoriasas
Other uses of this product should only be under the advice and supervision of a doctor.

***WARNING***
• For external use only
• avoid contact with the eyes
• if condition worsens, or symptoms persist for more than 7 days or clear up and occur again within a few days, stop use of this product and do not begin use of any other hydrocortisone product unless you have consulted a doctor
• do not use for the treatment of diaper rash.
• Keep out of reach of children.
• If swallowed, get medical help or consult a poison control center right away.

***DIRECTIONS***
***Adult and children (2 years and over):*** apply to affected not more than 3 to 4 times daily
***Children under 2 years:*** Consult a doctor.

***OTHER INFORMATION***
• store at room temperature 59-86°F (15-30°C) • do not freeze • do not use any opened or torn packets.

***INACTIVE INGREDIENTS***
emulsifying wax, ethanol, methylparaben, mineral oil, paraffin, petrolatum, propylparaben, purified water, white Wax.

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# Medi-Meclizine

## Drug Facts

**INGREDIENTS**
**Active ingredient (in each tablet):** Meclizine Hydrochloride 25mg **Purpose:** Antiemetic

**INACTIVE INGREDIENTS***
anhydrous lactose, colloidal silicon dioxide, corn starch, D&C yellow #10, magnesium stearate, microcrystalline cellulose, sodium starch glycolate

**USES**
For the prevention and treatment of nausea, vomiting, or dizziness associated with motion sickness. For the reduction of fever.

**WARNING**
***Do not use***
• for children under 12 years of age unless directed by a doctor • for frequent or prolonged use except under the advice of a doctor

***Ask a doctor before use if you have*** • breathing problems such as emphysema or chronic bronchitis • glaucoma
• difficulty in urination due to enlargement of the prostate gland if you are • taking sedatives or tranquilizers

***When using this product*** • drowsiness may occur • alcohol, sedatives and tranquilizers may increase the drowsiness effect
• avoid alcoholic beverages while taking this product • use caution when driving a motor vehicle or operating machinery

Do not exceed recommended dosage.
If pregnant or breast-feeding, ask a health professional before use.
Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away.

**DIRECTIONS**
• do not use more than directed • to prevent motion sickness, take the first dose one hour before starting activity
***Adults and children:*** (12 years and older) 1 to 2 tablets once daily or as directed by a doctor. Do not exceed 2 tablets in 24 hours.
***Children under 12 years:*** Do not give to children under 12 years of age.

**OTHER INFORMATION**
• store at room temperature 59- 86° F (15-30°C) • tamper-evident sealed packets • do not use any opened or torn packets

# Extra Strength APAP

## Drug Facts

**INGREDIENTS**
**Active ingredient (in each tablet):** Acetaminophen 500mg **Purpose:** Pain reliever/fever reducer

**INACTIVE INGREDIENTS***
corn starch, hypromellose, maltodextrin*, microcrystalline cellulose*, polyethylene glycol, povidone*, pregelatinized starch*, sodium starch glycolate*, stearic acid, titanium dioxide* *may contain

**USES**
For the temporary relief of minor aches and pains associated with
• headache • muscular aches • minor arthritis pain • common cold • toothache • menstrual cramps For the reduction of fever.

**WARNING**
**Liver warning:** This product contains acetaminophen. Severe liver damage may occur if you take:
• more than 8 tablets in 24 hours, which is the maximum daily amount
• with other drugs containing acetaminophen
• 3 or more alcoholic drinks every day while using this product

**Allergy alert:** Acetaminophen may cause severe skin reactions.
**Symptoms may include:**
• skin reddening • blisters • rash
If a skin reaction occurs, stop use and seek medical help right away.

***Do not use***
• with any other drug containing acetaminophen (prescription or nonprescription). If you are not sure whether a drug contains acetaminophen, ask a doctor or pharmacist.
• for more than 10 days for pain unless directed by a doctor
• for more than 3 days for fever unless directed by a doctor

***Ask a doctor before use if you have*** • liver disease

***Ask a doctor or pharmacist before use if*** • you are taking the blood thinning drug warfarin

***Stop using and ask a doctor if*** • symptoms do not improve • new symptoms occur • pain or fever persists or gets worse • redness or swelling is present

If pregnant or breast-feeding, ask a health professional before use. Keep out of reach of children. In case of accidental overdose, get medical help or contact a Poison Control Center right away. Prompt medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

**DIRECTIONS**
• do not use more than directed
***Adults and children:*** (12 years and older) Take 2 tablets every 4 to 6 hours as needed. Do not take more than 8 tablets in 24 hours. ***Children under 12 years:*** Do not give this adult strength product to children under 12 years of age; this will provide more than the recommended dose (overdose) and may cause liver damage.

**OTHER INFORMATION**
• store at room temperature 59°-86°F (15°-30°C) • tamper-evident sealed packets • do not use any opened or torn packets

# Diotame

## Drug Facts

**INGREDIENTS**
**Active ingredient (in each tablet):** Bismuth Subsalicylate 262mg (each tablet contains 102mg salicylate) **Purpose:** Upset stomach reliever/antidiarrheal

**INACTIVE INGREDIENTS***
acacia gum, aspartame, calcium carbonate, D&C red #27, dextrates, flavoring, magnesium stearate, maltodextrin, microcrystalline cellulose, silicon dioxide

**USES**

temporarily relieves • travelers' diarrhea • diarrhea
• upset stomach due to overindulgence in food and drink, including - heartburn - indigestion - nausea - gas - belching - fullness

**WARNING**
Reye's syndrome: Children and teenagers who have or are recovering from chicken pox or flu-like symptoms should not use this product. When using this product, if changes in behavior with nausea and vomiting occur, consult a doctor because these symptoms could be an early sign of Reye's Syndrome a rare but serious illness.

***Allergy alert: Contains salicylate. Do not take if you are:*** • allergic to salicylates (including aspirin) • taking other salicylate products

***Do not use if you have*** • bloody or black stool • an ulcer • a bleeding problem

***Ask a doctor before use if you have*** • fever • mucus in the stool

***Ask a doctor or pharmacist if you are taking any drug for*** • anticoagulation (thinning of the blood) • diabetes • gout • arthritis

***Stop use and ask a doctor if*** • symptoms get worse • ringing in the ears or loss of hearing occurs • diarrhea lasts more than 2 days

When using this product a temporary and harmless darkening of the tongue and/or stool may occur. Stool darkening should not be confused with melena.

If pregnant or breast-feeding, ask a health professional before use.
Keep this and all drugs out of reach of children. In case of accidental overdose, contact a physician or poison control center immediately.

**DIRECTIONS**
• do not use more than directed • chew or crush tablets completely before swallowing • do not swallow tablets whole
• use until diarrhea stops but not more than 2 days • drink plenty of clear fluids to help prevent dehydration, which may accompany diarrhea
• do not exceed 16 tablets in 24 hours
***Adults and children:*** (12 years and older) Chew 2 tablets every 1/2 to 1 hour or 4 tablets every hour as needed.
***Children under 12 years:*** Do not give to children under 12 years of age.

**OTHER INFORMATION**
• Phenylketonurics: contains phenylalanine 1.1mg per tablet • each tablet contains 73mg of elemental calcium • store at room temperature 59°-86°F (15°-30°C)
• tamper-evident sealed packets • do not use any opened or torn packets

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# Cold Relief

## Drug Facts

**INGREDIENTS**
**Active Ingredient (in each tablet):** Acetaminophen 325mg — **Purpose:** Pain reliever/fever reducer
**Active Ingredient (in each tablet):** Dextromethorphan Hydrobromide 15mg — **Purpose:** Cough suppressant
**Active Ingredient (in each tablet):** Guaifenesin 200mg — **Purpose:** Expectorant
**Active Ingredient (in each tablet):** Phenylephrine HCl 5mg — **Purpose:** Nasal decongestant

**INACTIVE INGREDIENTS***
maltodextrin, microcrystalline cellulose, povidone, sodium starch glycolate, starch, stearic acid

**USES**
Temporarily relieves these symptoms due to hay fever or other upper respiratory allergies
• cough • sore throat • minor aches and pains • headache • nasal congestion
• helps loosen phlegm (mucus) and thin bronchial secretions to make coughs more productive Temporarily reduces fever.

**WARNING**
***Liver warning:*** This product contains acetaminophen. Severe liver damage may occur if you take:
• more than 4,000mg of acetaminophen in 24 hours • with other drugs containing acetaminophen • 3 or more alcoholic drinks every day while using this product

***Allergy alert:*** Acetaminophen may cause severe skin reactions. Symptoms may include:
• skin reddening • blisters • rash If a skin reaction occurs, stop use and seek medical help right away.

***Do not use***
• with any other drug containing acetaminophen (prescription or nonprescription). If you are not sure whether a drug contains acetaminophen, ask a doctor or pharmacist.
• if you have ever had an allergic reaction to this product or any of the ingredients.
• if you are now taking a prescription monoamine oxidase inhibitor (MAOI) (certain drugs for depression, psychiatric or emotional conditions, or Parkinson's disease), or for 2 weeks after stopping the MAOI drug. If you do not know if your prescription drug contains an MAOI, ask a doctor or pharmacist before taking this product.

***Ask a doctor before use if you have***
• liver disease • heart disease • high blood pressure • thyroid disease • diabetes • trouble urinating due to an enlarged prostate gland
• cough that occurs with too much phlegm (mucus) • persistent or chronic cough that lasts as occurs with smoking, asthma, chronic bronchitis or emphysema

***Ask a doctor or pharmacist before use if you are*** • taking the blood thinning drug warfarin

***When using this product*** • do not use more than directed

***Stop use and ask a doctor if*** • new symptoms occur • redness or swelling is present • pain or nasal congestion gets worse or lasts for more than 7 days
• fever gets worse or lasts for more than 3 days • you get nervous, dizzy or sleepless • cough comes back or occurs with rash or headache that lasts.
These could be signs of a serious condition.

If pregnant or breast-feeding, ask a health professional before use.
Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away (1-800-222-1222).
Prompt medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

**DIRECTIONS**
***Adults and children:*** (12 years and older) Take 2 tablets with water every 6-8 hours as needed. Do not take more than 8 tablets in 24 hours.
***Children under 12 years:*** Do not give to children under 12 years of age.

**OTHER INFORMATION**
• store at room temperature 59°-86°F (15°-30°C) • avoid excessive heat and humidity • tamper-evident sealed packets • do not use any opened or torn packets

# Diphen

## Drug Facts

**INGREDIENTS**
**Active Ingredient (in each tablet):** Diphenhydramine HCl 25mg **Purpose:** Antihistamine

**INACTIVE INGREDIENTS***
carnauba wax*, colloidal silicon dioxide, croscarmellose sodium, D&C red #27, dicalcium phosphate*, hypromellose, lactose*,

magnesium stearate, microcrystalline cellulose, polyethylene glycol, polysorbate*, titanium dioxide *may contain

***USES***
Active ingredient (in each tablet): Diphenhydramine HCl 25mg Purpose: Antihistamine
***Inactive ingredients****
carnauba wax*, colloidal silicon dioxide, croscarmellose sodium, D&C red #27, dicalcium phosphate*, hypromellose, lactose*, magnesium stearate, microcrystalline cellulose, polyethylene glycol, polysorbate*, titanium dioxide * may contain

***WARNING***
***Do not use***
• to make a child sleepy • with any other product containing diphenhydramine, even one that is used on skin

***Ask a doctor before use if you have*** • a breathing problem such as emphysema or chronic bronchitis
• difficulty in urination due to enlargement of the prostate gland • glaucoma

***Ask a doctor or pharmacist before use if you are*** • taking sedatives or tranquilizers

***When using this product*** • marked drowsiness may occur • avoid alcohol beverages • alcohol, sedatives and tranquilizers may increase the drowsiness effect
• use caution when driving a motor vehicle or operating machinery • excitability may occur, especially in children

If pregnant or breast-feeding, ask a health professional before use.
Keep out of reach of children. In case of overdose, contact a physician or Poison Control Center immediately.

***DIRECTIONS***
• do not use more than directed
***Adults and children:*** (12 years and older) Take 1 to 2 caplets every 4 to 6 hours as needed. Do not take more than 12 caplets in 24 hours, or as directed by a doctor.
***Children under 12 years:*** Do not give to children under 12 years of age.

***OTHER INFORMATION***
• each caplet may contain: calcium 25mg • protect from light • use by expiration date on packet • store at room temperature 59°-86°F (15°-30°C)
• tamper-evident sealed packets • do not use any opened or torn packets

## Medi-Lyte

### *Drug Facts*

***INFORMATION***
Serving Size: 2 tablets
Servings Per Packet: 1

***OTHER INGREDIENTS***
microcrystalline cellulose, silicon dioxide, stearic acid, magnesium stearate

| ***Amount Per Serving*** | ***% Daily Value*** |
|---|---|
| Calcium (from 27.0 mg calcium carbonate) 10.8 mg | 1.06% |
| Potassium (from 80 mg potassium chloride) 40mg | 1.15% |
| Magnesium (from 20 mg magnesium oxide) 12 mg | 3.0% |
| Carbohydrates 6 mg | >1% |
| Calories 1.5 | >1% |
| Protein | 0% |
| Fat | 0% |

***USES***
Nutritional support for the following symptoms due to excessive loss of perspiration
• heat fatigue • muscle cramps • heat exhaustion • heat stroke • replaces lost electrolytes • helps provide rapid rehydration
* This statement has not been evaluated by the Food and Drug administration. This product is not intended to diagnose, treat, cure, or prevent any disease.

***WARNING***
If pregnant or breast-feeding, ask a health professional before use.
Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away.

***DIRECTIONS***
***Adults and children:*** (12 years and older) • take 2 tablets as needed with a full glass of water • may be repeated every hour as needed
• do not exceed 20 tablets in 24 hours ***Children under 12 years:*** • Do not give to children under 12 years of age

***OTHER INFORMATION***
• store at room temperature 59°-86°F (15°-30°C) • avoid excessive heat and humidity • tamper-evident sealed packets
• do not use any opened or torn packets • no sodium added

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## Alcalak

### *Drug Facts*

***INGREDIENTS***
**Active ingredient (in each tablet):** Calcium Carbonate 420mg **Purpose:** Antacid

***INACTIVE INGREDIENTS****
aspartame*, croscarmellose sodium*, gum acacia*, magnesium stearate, maltodextrin, mineral oil*, mint flavor, sorbitol*, sucrose* *may contain

***USES***
For the relief of the following symptoms associated with • acid indigestion • sour stomach • heartburn • upset stomach

***WARNING***
***Do not use***
• the maximum dosage of this product for more than 2 weeks, except under the advice and supervision of a physician, or take more than 19 tablets in a 24 hour period.

***Ask a doctor or pharmacist before use if you are*** • presently taking a prescription drug. Antacids may interact with certain prescription drugs.

***Stop use and ask a doctor if*** • symptoms last more than 2 weeks

If pregnant or breastfeeding, ask a health professional before use.
Keep out of reach of children.

***DIRECTIONS***
• do not use more than directed
***Adults and children:*** (12 years and older) Chew 2 tablets every 2 or 3 hours as symptoms occur or as directed by a physician. Do not exceed 19 tablets in 24 hours.
***Children under 12 years:*** Do not give to children under 12 years of age.

***OTHER INFORMATION***

• Phenylketonurics: contains phenylalanine 1.5mg per tablet • each tablet contains 168mg of elemental calcium
• store at room temperature 59°-86°F (15°-30°C) in a dry place • tamper-evident sealed packets • do not use any opened or torn packets

# I-Prin

## Drug Facts

***INGREDIENTS***
**Active ingredient (in each tablet):** Ibuprofen (NSAID*) 200mg **Purpose:** Pain reliever/fever reducer *nonsteroidal anti-inflammatory drug

***INACTIVE INGREDIENTS****
carnauba wax*, corn starch, hypromellose*, iron oxide red, lactose*, magnesium stearate*, microcrystalline cellulose*, polydextrose*, polyethylene glycol, polyvinyl alcohol*, povidone (K-30)*, silicon dioxide, sodium starch glycolate, stearic acid, talc*, titanium dioxide *may contain

***USES***
Temporarily relieves minor aches and pains associated with
• headache • toothache • backache • menstrual cramps • common cold • muscular aches • minor arthritis pain Temporarily reduces fever.

***WARNING***
**Allergy alert:** Ibuprofen may cause a severe allergic reaction, especially in people allergic to aspirin.
**Symptoms may include:** • hives • skin reddening • facial swelling • rash • asthma (wheezing) • blisters • shock
If an allergic reaction occurs, stop use and seek medical help right away.

**Stomach bleeding warning:** This product contains an NSAID, which may cause severe stomach bleeding.
**The chance is higher if you:** • are age 60 or older • have had stomach ulcers or bleeding problems • take a blood thinning (anticoagulant) or steroid drug
• take other drugs containing prescription or nonprescription NSAIDs (aspirin, ibuprofen, naproxen, or others)
• have 3 or more alcoholic drinks every day while using this product • take more or for a longer time than directed

Heart attack or stroke warning: NSAIDS, except aspirin, increase the risk of heart attack, heart failure, and stroke.
These can be fatal. The risk is higher if you use more than directed or for longer than directed.

***Do not use***
• if you have ever had an allergic reaction to any other pain reliever/ fever reducer • right before or after heart surgery

***Ask a doctor before use if*** • you have problems or serious side effects from taking pain relievers or fever reducers
• stomach bleeding warning applies to you • you have a history of stomach problems such as heartburn
• you have high blood pressure, heart disease, liver cirrhosis, kidney disease, asthma or had a stroke • you are taking a diuretic

***Ask a doctor or pharmacist before use if you are*** • taking aspirin for heart attack or stroke, because ibuprofen may decrease this benefit of aspirin
• under a doctor's care for any serious condition • taking any other drug

***When using this product*** • take with food or milk if stomach upset occurs

***Stop use and ask a doctor if***
• you experience any of the following signs of stomach bleeding: - feel faint - vomit blood - have bloody or black stools - have stomach pain that does not get better
• you have symptoms of heart problems or stroke - chest pain - trouble breathing - weakness in one part or side of body - slurred speech - leg swelling
• pain gets worse or lasts more than 10 days • fever gets worse or lasts more than 3 days
• redness or swelling is present in the painful area • any new or unexpected symptoms occur

If pregnant or breast-feeding, ask a health professional before use. It is especially important not to use ibuprofen during the last 3 months of pregnancy unless specifically directed to do so by a doctor because it may cause problems in the unborn child or complications during delivery.
Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away.

***DIRECTIONS***
• do not take more than directed • the smallest effective dose should be used • do not take longer than 10 days, unless directed by a doctor (see Warnings)
***Adults and children:*** (12 years and older) Take 1 tablet every 4 to 6 hours while symptoms persist. If pain or fever does not respond to 1 tablet, 2 tablets may be used.
Do not exceed 6 tablets in 24 hours, unless directed by a doctor. ***Children under 12 years:*** Do not give to children under 12 years of age.

***OTHER INFORMATION***
• read all product information before using • store at 68-77°F (20-25°C) • avoid excessive heat 104°F (above 40°C)
• tamper-evident sealed packets • do not use any opened or torn packets

# Loradamed

## Drug Facts

***INGREDIENTS***
**Active ingredient (in each tablet):** Loratadine 10mg **Purpose:** Antihistamine

***INACTIVE INGREDIENTS****
corn starch, lactose monohydrate, magnesium stearate, pregelatinized starch

***USES***
Temporarily relieves these symptoms due to hay fever or other upper respiratory allergies
• runny nose • itchy, watery eyes • sneezing • itching of the nose or throat

***WARNING***
Do not use if you have ever had an allergic reaction to this product or any of its ingredients.

Ask a doctor before use if you have liver or kidney disease. Your doctor should determine if you need a different dose.

When using this product do not take more than directed. Taking more than directed may cause drowsiness.

***Stop use and ask a doctor if*** • *an allergic reaction to this product occurs. Seek medical help right away.*

If pregnant or breast-feeding, ask a health professional before use.
Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

***DIRECTIONS***
***Adults and children:*** (12 years and older) Take 1 tablet daily; not more than 1 tablet in 24 hours. **Children under 12 years:** Do not give to children under 12 years of age.
***Consumers with liver or kidney disease:*** Ask a doctor before using.

***OTHER INFORMATION***
• store at room temperature 68°-77°F (20°-25°C) • protect from excessive moisture • tamper-evident sealed packets • do not use any opened or torn packets

# DIPHEN

benzocaine, benzalkonium chloride, lidocaine hydrochloride, hydrocortisone, bacitracin zinc, neomycin sulfate, polymyxin b sulfate, calcium carbonate, ibuprofen, loratadine, acetaminophen, dextromethorphan hydrobromide, guaifenesin, phenylephrine hydrochloride, diphenhydramine hydrochloride, potassium chloride, magnesium oxide, meclizine hydrochloride, and bismuth subsalicylate kit

| Product Information | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:82652-021 |

| Packaging | | | | |
|---|---|---|---|---|
| **#** | **Item Code** | **Package Description** | **Marketing Start Date** | **Marketing End Date** |
| **1** | NDC:82652-021-01 | 1 in 1 CARTON | 05/10/2022 | |

| Quantity of Parts | | |
|---|---|---|
| **Part #** | **Package Quantity** | **Total Product Quantity** |
| **Part 1** | 4 PACKET | 3 g |
| **Part 2** | 2 PACKET | 1.8 g |
| **Part 3** | 4 PACKET | 3.6 g |
| **Part 4** | 8 PACKET | 4 g |
| **Part 5** | 2 PACKET | 4 |
| **Part 6** | 4 PACKET | 8 |
| **Part 7** | 4 PACKET | 4 |
| **Part 8** | 4 PACKET | 8 |
| **Part 9** | 4 PACKET | 4 |
| **Part 10** | 2 PACKET | 4 |
| **Part 11** | 3 PACKET | 6 |
| **Part 12** | 4 PACKET | 8 |
| **Part 13** | 3 PACKET | 6 |

## Part 1 of 13

## PAIN RELIEF

benzocaine liquid

| Product Information | |
|---|---|
| **Item Code (Source)** | NDC:82652-033(NDC:61010-8100) |
| **Route of Administration** | TOPICAL |

| Active Ingredient/Active Moiety | | |
|---|---|---|
| **Ingredient Name** | **Basis of Strength** | **Strength** |

| | | |
|---|---|---|
| **benzocaine** (UNII: U3RSY48JW5) (benzocaine - UNII:U3RSY48JW5) | benzocaine | 200 mg in 1 g |

### Inactive Ingredients

| Ingredient Name | Strength |
|---|---|
| **polyethylene glycol 400** (UNII: B697894SGQ) | |
| **polyethylene glycol 3350** (UNII: G2M7P15E5P) | |
| **peppermint oil** (UNII: AV092KU4JH) | |
| **saccharin sodium** (UNII: SB8ZUX40TY) | |
| **sorbic acid** (UNII: X045WJ989B) | |

### Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| **1** | NDC:82652-033-01 | 0.75 g in 1 PACKET; Type 0: Not a Combination Product | | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| OTC monograph not final | part356 | 05/01/2010 | |

## Part 2 of 13

## BURN

benzalkonium chloride and lidocaine hydrochloride cream

### Product Information

| | |
|---|---|
| **Item Code (Source)** | NDC:82652-029(NDC:47682-940) |
| **Route of Administration** | TOPICAL |

### Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|---|---|---|
| **Benzalkonium Chloride** (UNII: F5UM2KM3W7) (Benzalkonium - UNII:7N6JUD5X6Y) | Benzalkonium Chloride | 1.3 mg in 1 g |
| **Lidocaine Hydrochloride** (UNII: V13007Z41A) (Lidocaine - UNII:98PI200987) | Lidocaine Hydrochloride Anhydrous | 5 mg in 1 g |

### Inactive Ingredients

| Ingredient Name | Strength |
|---|---|
| **Alcohol** (UNII: 3K9958V90M) | |

| | |
|---|---|
| **Methylparaben** (UNII: A2I8C7HI9T) | |
| **Mineral Oil** (UNII: T5L8T28FGP) | |
| **Paraffin** (UNII: I9O0E3H2ZE) | |
| **Propylparaben** (UNII: Z8IX2SC1OH) | |
| **Water** (UNII: 059QF0KO0R) | |
| **White Petrolatum** (UNII: B6E5W8RQJ4) | |
| **White Wax** (UNII: 7G1J5DA97F) | |

## Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| **1** | NDC:82652-029-01 | 0.9 g in 1 PACKET; Type 0: Not a Combination Product | | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| OTC MONOGRAPH NOT FINAL | part348 | 08/29/2017 | |

## Part 3 of 13

## HYDROCORTISONE

hydrocortisone cream

## Product Information

| | |
|---|---|
| **Item Code (Source)** | NDC:82652-027(NDC:47682-923) |
| **Route of Administration** | TOPICAL |

## Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|---|---|---|
| **HYDROCORTISONE ACETATE** (UNII: 3X7931PO74) (Hydrocortisone - UNII:W4X0X7BPJ) | HYDROCORTISONE ACETATE | 10 mg in 1 g |

## Inactive Ingredients

| Ingredient Name | Strength |
|---|---|
| **Alcohol** (UNII: 3K9958V90M) | |
| **Methylparaben** (UNII: A2I8C7HI9T) | |
| **Mineral Oil** (UNII: T5L8T28FGP) | |
| **Paraffin** (UNII: I9O0E3H2ZE) | |
| **Petrolatum** (UNII: 4T6H12BN9U) | |
| **Propylparaben** (UNII: Z8IX2SC1OH) | |

| | |
|---|---|
| **Water** (UNII: 059QF0KO0R) | |
| **White Wax** (UNII: 7G1J5DA97F) | |

## Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| **1** | NDC:82652-027-01 | 0.9 g in 1 PACKET; Type 0: Not a Combination Product | | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| OTC MONOGRAPH NOT FINAL | part348 | 06/01/2021 | |

# Part 4 of 13

# TRIPLE ANTIBIOTIC

bacitracin zinc, neomycin sulfate, and polymyxin b sulfate ointment

## Product Information

| | |
|---|---|
| **Item Code (Source)** | NDC:82652-028(NDC:47682-932) |
| **Route of Administration** | TOPICAL |

## Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|---|---|---|
| **Bacitracin Zinc** (UNII: 89Y4M234ES) (Bacitracin - UNII:58H6RWO52I) | Bacitracin | 400 [USP'U] in 1 g |
| **Neomycin Sulfate** (UNII: 057Y626693) (Neomycin - UNII:I16QD7X297) | Neomycin | 3.5 mg in 1 g |
| **Polymyxin B Sulfate** (UNII: 19371312D4) (Polymyxin B - UNII:J2VZ07J96K) | Polymyxin B | 5000 [USP'U] in 1 g |

## Inactive Ingredients

| Ingredient Name | Strength |
|---|---|
| **Polyethylene Glycol 3350** (UNII: G2M7P15E5P) | |
| **Polyethylene Glycol 400** (UNII: B697894SGQ) | |
| **Saccharin Sodium** (UNII: SB8ZUX40TY) | |
| **FD&C Blue No. 1** (UNII: H3R47K3TBD) | |
| **FD&C Green No. 3** (UNII: 3P3ONR6O1S) | |
| **D&C Green No. 5** (UNII: 8J6RDU8L9X) | |
| **FD&C Red No. 3** (UNII: PN2ZH5LOQY) | |
| **D&C Red No. 28** (UNII: 767IP0Y5NH) | |
| **FD&C Red No. 40** (UNII: WZB9127XOA) | |

| | |
|---|---|
| **FD&C Yellow No. 5** (UNII: I753WB2F1M) | |
| **FD&C Yellow No. 6** (UNII: H77VEI93A8) | |

### Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| **1** | NDC:82652-028-01 | 0.5 g in 1 PACKET; Type 0: Not a Combination Product | | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| OTC MONOGRAPH FINAL | part333B | 06/04/2018 | |

## Part 5 of 13

## ALCALAK

calcium carbonate tablet, chewable

### Product Information

| | |
|---|---|
| **Item Code (Source)** | NDC:82652-023(NDC:47682-201) |
| **Route of Administration** | ORAL |

### Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|---|---|---|
| **Calcium Carbonate** (UNII: H0G9379FGK) (CALCIUM CATION - UNII:2M83C4R6ZB, CARBONATE ION - UNII:7UJQ5OPE7D) | Calcium Carbonate | 420 mg |

### Inactive Ingredients

| Ingredient Name | Strength |
|---|---|
| **Aspartame** (UNII: Z0H242BBR1) | |
| **Croscarmellose Sodium** (UNII: M28OL1HH48) | |
| **Acacia** (UNII: 5C5403N26O) | |
| **Magnesium Stearate** (UNII: 70097M6I30) | |
| **Maltodextrin** (UNII: 7CVR7L4A2D) | |
| **Mineral Oil** (UNII: T5L8T28FGP) | |
| **Sorbitol** (UNII: 506T60A25R) | |
| **Sucrose** (UNII: C151H8M554) | |

| Product Characteristics | | | |
|---|---|---|---|
| **Color** | WHITE | **Score** | no score |
| **Shape** | ROUND | **Size** | 11mm |
| **Flavor** | | **Imprint Code** | AZ;036 |
| **Contains** | | | |

| Packaging | | | | |
|---|---|---|---|---|
| **#** | **Item Code** | **Package Description** | **Marketing Start Date** | **Marketing End Date** |
| **1** | NDC:82652-023-01 | 2 in 1 PACKET; Type 0: Not a Combination Product | | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| OTC MONOGRAPH FINAL | part331 | 06/15/2014 | |

## Part 6 of 13

## I-PRIN

ibuprofen tablet, film coated

| Product Information | |
|---|---|
| **Item Code (Source)** | NDC:82652-024(NDC:47682-683) |
| **Route of Administration** | ORAL |

| Active Ingredient/Active Moiety | | |
|---|---|---|
| **Ingredient Name** | **Basis of Strength** | **Strength** |
| **Ibuprofen** (UNII: WK2XYI10QM) (Ibuprofen - UNII:WK2XYI10QM) | Ibuprofen | 200 mg |

| Inactive Ingredients | |
|---|---|
| **Ingredient Name** | **Strength** |
| **Carnauba Wax** (UNII: R12CBM0EIZ) | |
| **Starch, Corn** (UNII: O8232NY3SJ) | |
| **Hypromellose, Unspecified** (UNII: 3NXW29V3WO) | |
| **Ferric Oxide Red** (UNII: 1K09F3G675) | |
| **Lactose, Unspecified Form** (UNII: J2B2A4N98G) | |
| **Magnesium Stearate** (UNII: 70097M6I30) | |
| **Microcrystalline Cellulose** (UNII: OP1R32D61U) | |
| **Polydextrose** (UNII: VH2XOU12IE) | |

| | |
|---|---|
| **Polyethylene Glycol, Unspecified** (UNII: 3WQ0SDW1A) | |
| **Polyvinyl Alcohol, Unspecified** (UNII: 532B59J990) | |
| **Povidone K30** (UNII: U725QWY32X) | |
| **Silicon Dioxide** (UNII: ETJ7Z6XBU4) | |
| **SODIUM STARCH GLYCOLATE TYPE A** (UNII: H8AV0SQX4D) | |
| **Stearic Acid** (UNII: 4ELV7Z65AP) | |
| **Talc** (UNII: 7SEV7J4R1U) | |
| **Titanium Dioxide** (UNII: 15FIX9V2JP) | |

### Product Characteristics

| | | | |
|---|---|---|---|
| **Color** | BROWN | **Score** | no score |
| **Shape** | ROUND | **Size** | 10mm |
| **Flavor** | | **Imprint Code** | 44;291 |
| **Contains** | | | |

### Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| **1** | NDC:82652-024-01 | 2 in 1 PACKET; Type 0: Not a Combination Product | | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| ANDA | ANDA075010 | 02/01/2021 | |

## Part 7 of 13

## LORADAMED

loratadine tablet, film coated

### Product Information

| | |
|---|---|
| **Item Code (Source)** | NDC:82652-022(NDC:47682-203) |
| **Route of Administration** | ORAL |

### Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|---|---|---|
| **Loratadine** (UNII: 7AJO3BO7QN) (Loratadine - UNII:7AJO3BO7QN) | Loratadine | 10 mg |

### Inactive Ingredients

| Ingredient Name | Strength |
|---|---|
| **Starch, Corn** (UNII: O8232NY3SJ) | |
| **Lactose Monohydrate** (UNII: EWQ57Q8I5X) | |
| **Magnesium Stearate** (UNII: 70097M6I30) | |

| Product Characteristics | | | |
|---|---|---|---|
| **Color** | WHITE | **Score** | no score |
| **Shape** | ROUND | **Size** | 6mm |
| **Flavor** | | **Imprint Code** | RX;526 |
| **Contains** | | | |

**Packaging**

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| **1** | NDC:82652-022-01 | 1 in 1 PACKET; Type 0: Not a Combination Product | | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| ANDA | ANDA076134 | 12/30/2008 | |

## Part 8 of 13

## COLD RELIEF

acetaminophen, dextromethorphan hydrobromide, guaifenesin, and phenylephrine hydrochloride tablet

| Product Information | |
|---|---|
| **Item Code (Source)** | NDC:82652-032(NDC:47682-725) |
| **Route of Administration** | ORAL |

**Active Ingredient/Active Moiety**

| Ingredient Name | Basis of Strength | Strength |
|---|---|---|
| **Acetaminophen** (UNII: 362O9ITL9D) (Acetaminophen - UNII:362O9ITL9D) | Acetaminophen | 325 mg |
| **Dextromethorphan Hydrobromide** (UNII: 9D2RTI9KYH) (Dextromethorphan - UNII:7355X3ROTS) | Dextromethorphan Hydrobromide | 15 mg |
| **Guaifenesin** (UNII: 495W7451VQ) (Guaifenesin - UNII:495W7451VQ) | Guaifenesin | 200 mg |
| **Phenylephrine Hydrochloride** (UNII: 04JA59TNSJ) (Phenylephrine - UNII:1WS297W6MV) | Phenylephrine Hydrochloride | 5 mg |

## Inactive Ingredients

| Ingredient Name | Strength |
|---|---|
| **Maltodextrin** (UNII: 7CVR7L4A2D) | |
| **Microcrystalline Cellulose** (UNII: OP1R32D61U) | |
| **Povidone, Unspecified** (UNII: FZ989GH94E) | |
| **SODIUM STARCH GLYCOLATE TYPE A** (UNII: H8AV0SQX4D) | |
| **Stearic Acid** (UNII: 4ELV7Z65AP) | |

## Product Characteristics

| | | | |
|---|---|---|---|
| **Color** | WHITE | **Score** | no score |
| **Shape** | ROUND | **Size** | 12mm |
| **Flavor** | | **Imprint Code** | FR;12 |
| **Contains** | | | |

## Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| **1** | NDC:82652-032-01 | 2 in 1 PACKET; Type 0: Not a Combination Product | | |

# Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| OTC MONOGRAPH FINAL | part341 | 02/01/2021 | |

# Part 9 of 13

# DIPHEN

diphenhydramine hydrochloride tablet, film coated

## Product Information

| | |
|---|---|
| **Item Code (Source)** | NDC:82652-031(NDC:47682-166) |
| **Route of Administration** | ORAL |

## Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|---|---|---|
| **Diphenhydramine Hydrochloride** (UNII: TC2D6JAD40) (Diphenhydramine - UNII:8GTS82S83M) | Diphenhydramine Hydrochloride | 25 mg |

## Inactive Ingredients

| Ingredient Name | Strength |
|---|---|
| **Carnauba Wax** (UNII: R12CBM0EIZ) | |
| **Silicon Dioxide** (UNII: ETJ7Z6XBU4) | |
| **Croscarmellose Sodium** (UNII: M28OL1HH48) | |
| **D&C Red No. 27** (UNII: 2LRS185U6K) | |
| **Anhydrous Dibasic Calcium Phosphate** (UNII: L11K75P92J) | |
| **Hypromellose, Unspecified** (UNII: 3NXW29V3WO) | |
| **Lactose, Unspecified Form** (UNII: J2B2A4N98G) | |
| **Magnesium Stearate** (UNII: 70097M6I30) | |
| **Microcrystalline Cellulose** (UNII: OP1R32D61U) | |
| **Polyethylene Glycol, Unspecified** (UNII: 3WQ0SDW1A) | |
| **Titanium Dioxide** (UNII: 15FIX9V2JP) | |

## Product Characteristics

| | | | |
|---|---|---|---|
| **Color** | PINK | **Score** | no score |
| **Shape** | CAPSULE | **Size** | 11mm |
| **Flavor** | | **Imprint Code** | 048;D |
| **Contains** | | | |

## Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| **1** | NDC:82652-031-01 | 1 in 1 PACKET; Type 0: Not a Combination Product | | |

# Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| OTC MONOGRAPH FINAL | part341 | 02/01/2021 | |

# Part 10 of 13

# MEDI-LYTE

calcium carbonate, potassium chloride, and magnesium oxide tablet

## Product Information

| | |
|---|---|
| **Route of Administration** | ORAL |

## Active Ingredient/Active Moiety

| | Basis of |
|---|---|

| Ingredient Name | Basis of Strength | Strength |
|---|---|---|
| **Calcium Carbonate** (UNII: H0G9379FGK) (Calcium Cation - UNII:2M83C4R6ZB, Carbonate Ion - UNII:7UJQ5OPE7D) | Calcium Carbonate | 27 mg |
| **Potassium Chloride** (UNII: 660YQ98I10) (POTASSIUM CATION - UNII:295O53K152, CHLORIDE ION - UNII:Q32ZN48698) | Potassium Chloride | 80 mg |
| **Magnesium Oxide** (UNII: 3A3U0GI71G) (MAGNESIUM CATION - UNII:T6V3LHY838) | Magnesium Oxide | 20 mg |

## Inactive Ingredients

| Ingredient Name | Strength |
|---|---|
| **Microcrystalline Cellulose** (UNII: OP1R32D61U) | |
| **Silicon Dioxide** (UNII: ETJ7Z6XBU4) | |
| **Stearic Acid** (UNII: 4ELV7Z65AP) | |
| **Magnesium Stearate** (UNII: 70097M6I30) | |

## Product Characteristics

| | | | |
|---|---|---|---|
| **Color** | WHITE | **Score** | no score |
| **Shape** | ROUND | **Size** | 9mm |
| **Flavor** | | **Imprint Code** | |
| **Contains** | | | |

## Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| **1** | | 2 in 1 PACKET; Type 0: Not a Combination Product | | |

# Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| DIETARY SUPPLEMENT | | 05/10/2022 | |

# Part 11 of 13

# MEDI-MECLIZINE

meclizine hydrochloride tablet

## Product Information

| | |
|---|---|
| **Item Code (Source)** | NDC:82652-030(NDC:47682-481) |
| **Route of Administration** | ORAL |

| Active Ingredient/Active Moiety | | |
|---|---|---|
| **Ingredient Name** | **Basis of Strength** | **Strength** |
| **Meclizine Hydrochloride** (UNII: HDP7W44CIO) (Meclizine - UNII:3L5TQ84570) | Meclizine Hydrochloride | 25 mg |

| Inactive Ingredients | |
|---|---|
| **Ingredient Name** | **Strength** |
| **Anhydrous Lactose** (UNII: 3SY5LH9PMK) | |
| **Silicon Dioxide** (UNII: ETJ7Z6XBU4) | |
| **Starch, Corn** (UNII: O8232NY3SJ) | |
| **D&C Yellow No. 10** (UNII: 35SW5USQ3G) | |
| **Magnesium Stearate** (UNII: 70097M6I30) | |
| **Microcrystalline Cellulose** (UNII: OP1R32D61U) | |
| **SODIUM STARCH GLYCOLATE TYPE A** (UNII: H8AV0SQX4D) | |

| Product Characteristics | | | |
|---|---|---|---|
| **Color** | YELLOW | **Score** | no score |
| **Shape** | ROUND | **Size** | 9mm |
| **Flavor** | | **Imprint Code** | 44;403 |
| **Contains** | | | |

| Packaging | | | | |
|---|---|---|---|---|
| **#** | **Item Code** | **Package Description** | **Marketing Start Date** | **Marketing End Date** |
| **1** | NDC:82652-030-01 | 2 in 1 PACKET; Type 0: Not a Combination Product | | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| OTC MONOGRAPH FINAL | part336 | 12/14/2020 | |

## Part 12 of 13

## EXTRA STRENGTH APAP

acetaminophen tablet, film coated

| Product Information | |
|---|---|
| **Item Code (Source)** | NDC:82652-025(NDC:47682-043) |
| **Route of Administration** | ORAL |

| Active Ingredient/Active Moiety | | |
|---|---|---|
| **Ingredient Name** | **Basis of Strength** | **Strength** |
| **Acetaminophen** (UNII: 362O9ITL9D) (Acetaminophen - UNII:362O9ITL9D) | Acetaminophen | 500 mg |

| Inactive Ingredients | |
|---|---|
| **Ingredient Name** | **Strength** |
| **Starch, Corn** (UNII: O8232NY3SJ) | |
| **Hypromellose, Unspecified** (UNII: 3NXW29V3WO) | |
| **Maltodextrin** (UNII: 7CVR7L4A2D) | |
| **Microcrystalline Cellulose** (UNII: OP1R32D61U) | |
| **Polyethylene Glycol, Unspecified** (UNII: 3WQ0SDW1A) | |
| **SODIUM STARCH GLYCOLATE TYPE A** (UNII: H8AV0SQX4D) | |
| **Stearic Acid** (UNII: 4ELV7Z65AP) | |
| **Titanium Dioxide** (UNII: 15FIX9V2JP) | |

| Product Characteristics | | | |
|---|---|---|---|
| **Color** | WHITE | **Score** | no score |
| **Shape** | ROUND | **Size** | 12mm |
| **Flavor** | | **Imprint Code** | FR;33 |
| **Contains** | | | |

| Packaging | | | | |
|---|---|---|---|---|
| **#** | **Item Code** | **Package Description** | **Marketing Start Date** | **Marketing End Date** |
| **1** | NDC:82652-025-01 | 2 in 1 PACKET; Type 0: Not a Combination Product | | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| OTC MONOGRAPH NOT FINAL | part343 | 02/01/2021 | |

## Part 13 of 13

## DIOTAME

bismuth subsalicylate tablet, chewable

| Product Information | |
|---|---|
| **Item Code (Source)** | NDC:82652-026(NDC:47682-210) |
| **Route of Administration** | ORAL |

## Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|---|---|---|
| **Bismuth Subsalicylate** (UNII: 62TEY51RR1) (SALICYLIC ACID - UNII:O414PZ4LPZ, BISMUTH CATION - UNII:ZS9CD1I8YE) | Bismuth Subsalicylate | 262 mg |

## Inactive Ingredients

| Ingredient Name | Strength |
|---|---|
| **Acacia** (UNII: 5C5403N26O) | |
| **Aspartame** (UNII: Z0H242BBR1) | |
| **Calcium Carbonate** (UNII: H0G9379FGK) | |
| **D&C Red No. 27** (UNII: 2LRS185U6K) | |
| **Dextrates** (UNII: G263MI44RU) | |
| **Magnesium Stearate** (UNII: 70097M6I30) | |
| **Maltodextrin** (UNII: 7CVR7L4A2D) | |
| **Microcrystalline Cellulose** (UNII: OP1R32D61U) | |
| **Silicon Dioxide** (UNII: ETJ7Z6XBU4) | |

## Product Characteristics

| | | | |
|---|---|---|---|
| **Color** | PINK | **Score** | no score |
| **Shape** | ROUND | **Size** | 16mm |
| **Flavor** | PEPPERMINT | **Imprint Code** | RH;046 |
| **Contains** | | | |

## Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| **1** | NDC:82652-026-01 | 2 in 1 PACKET; Type 0: Not a Combination Product | | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| OTC MONOGRAPH FINAL | part335 | 04/01/2014 | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| ANDA | ANDA076134 | 05/10/2022 | |

## Labeler - Remedy Pack LLC (101454060)

## Establishment

| Name | Address | ID/FEI | Business Operations |
|---|---|---|---|
| Safetec of America, Inc | | 874965262 | MANUFACTURE(82652-021) |

## Establishment

| Name | Address | ID/FEI | Business Operations |
|---|---|---|---|
| Remedy Pack LLC | | 101454060 | RELABEL(82652-033, 82652-029, 82652-027, 82652-028, 82652-023, 82652-024, 82652-022, 82652-032, 82652-031, 82652-030, 82652-025, 82652-026) , REPACK(82652-033, 82652-029, 82652-027, 82652-028, 82652-023, 82652-024, 82652-022, 82652-032, 82652-031, 82652-030, 82652-025, 82652-026) |

## Establishment

| Name | Address | ID/FEI | Business Operations |
|---|---|---|---|
| Allegiant Health | | 079501930 | MANUFACTURE(82652-021) |

## Establishment

| Name | Address | ID/FEI | Business Operations |
|---|---|---|---|
| LNK International Inc. | | 966812120 | MANUFACTURE(82652-021) |

## Establishment

| Name | Address | ID/FEI | Business Operations |
|---|---|---|---|
| Granules India Limited | | 918609236 | MANUFACTURE(82652-021) |

## Establishment

| Name | Address | ID/FEI | Business Operations |
|---|---|---|---|
| Ohm Laboratories | | 051565745 | MANUFACTURE(82652-021) |

Revised: 7/2022 Remedy Pack LLC